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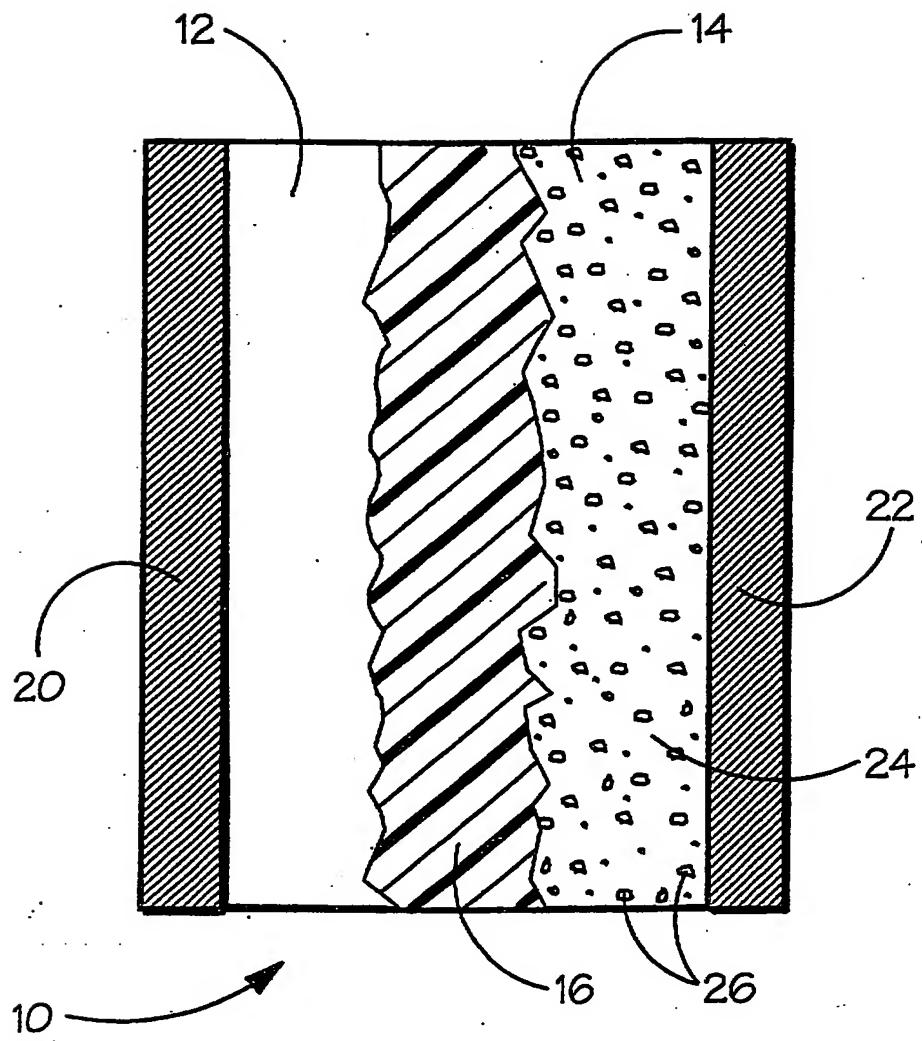
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(54) Abstract Title
Lithium metal dispersion in secondary battery anodes

(57) The present invention is a secondary battery having a high specific capacity and good cycleability, and that can be used safely. The secondary battery is manufactured to include an anode formed from a host material capable of absorbing and desorbing lithium in an electrochemical system such as a carbonaceous material, and lithium metal dispersed in the host material. The freshly prepared anodes of the invention are combined with a positive electrode including an active material, a separator that separates the positive electrode and the negative electrode, and an electrolyte in communication with the positive electrode and the negative electrode. The present invention also includes a method of preparing a freshly prepared anode and a method of operating a secondary battery including the anode of the invention.

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LITHIUM METAL DISPERSION IN SECONDARY BATTERY ANODES

Field of the Invention

The present invention relates to secondary batteries having high specific capacities and particularly to anodes for secondary batteries comprising a host material such as a carbonaceous material capable of absorbing and desorbing lithium in an electrochemical system and lithium metal dispersed in the host material.

Background of the Invention

Lithium and lithium-ion secondary or rechargeable batteries have recently found use in certain applications such as in cellular phones, camcorders, and laptop computers, and even more recently, in larger power applications such as in electric vehicles and hybrid electric vehicles. It is preferred in these applications that the secondary batteries have the highest specific capacity possible but still provide safe operating conditions and good cycleability so that the high specific capacity is maintained in subsequent recharging and discharging cycles.

Although there are various constructions for secondary batteries, each construction includes a positive electrode (or cathode), a negative electrode (or anode), a separator that separates the cathode and anode, and an electrolyte in electrochemical communication with the cathode and anode. For secondary lithium batteries, lithium ions are transferred from the anode to the cathode through the electrolyte when the secondary battery is being discharged, i.e., used for its specific application. During this process, electrons are collected from the anode and pass to the cathode through an external circuit. When the secondary battery is being charged or recharged, the lithium ions are transferred from the cathode to the anode through the electrolyte.

Historically, secondary lithium batteries were produced using non-lithiated compounds having high specific capacities such as TiS_2 , MoS_2 , MnO_2 and V_2O_5 , as the cathode active materials. These cathode active materials were coupled with a lithium metal anode. When the secondary battery was discharged, lithium ions were transferred from the lithium metal anode to the cathode through the electrolyte. Unfortunately, upon cycling, the lithium metal developed dendrites that ultimately caused unsafe conditions in the battery. As a result, the production of these types of secondary batteries was stopped in the early 1990's in favor of lithium-ion batteries.

Lithium-ion batteries typically use lithium metal oxides such as

10 $LiCoO_2$ and $LiNiO_2$ as cathode active materials coupled with a carbon-based anode. In these batteries, the lithium dendrite formation on the anode is avoided thereby making the battery safer. However, the lithium, the amount of which determines the battery capacity, is totally supplied from the cathode. This limits the choice of cathode active materials because the active materials must contain removable lithium.

15 Furthermore, the delithiated products corresponding to $LiCoO_2$ and $LiNiO_2$ that are formed during charging (e.g. Li_xCoO_2 and Li_xNiO_2 where $0.4 < x < 1.0$) and overcharging (i.e. Li_xCoO_2 and Li_xNiO_2 where $x < 0.4$) are not stable. In particular, these delithiated products tend to react with the electrolyte and generate heat, which raises safety concerns.

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Summary of the Invention

The present invention is a secondary battery having a high specific capacity and good cycleability and that operates safely. In accordance with the invention, the freshly prepared, secondary battery includes an anode that is formed of a host material capable of absorbing and desorbing lithium in an electrochemical system and lithium metal dispersed in the host material. Preferably, the lithium metal is a finely divided lithium powder and more preferably has a mean particle size of less than about 20 microns. The host material comprises one or more materials selected from the group consisting of carbonaceous materials, Si, Sn, tin oxides, composite tin alloys, transition metal oxides, lithium metal nitrides and lithium metal oxides.

Preferably, the host material comprises a carbonaceous material and more preferably comprises graphite.

The freshly prepared, secondary batteries of the invention include a positive electrode including an active material, a negative electrode comprising a host material capable of absorbing and desorbing lithium in an electrochemical system and lithium metal dispersed in the host material, a separator separating the positive electrode and the negative electrode and an electrolyte in communication with the positive electrode and the negative electrode. Preferably, the cathode active material is a compound that can be lithiated at an electrochemical potential of 2.0 to 5.0 V versus lithium. For example, the cathode active material can be MnO_2 , V_2O_5 or MoS_2 , or a mixture thereof. The lithium metal in the anode is preferably a finely divided lithium powder and more preferably has a mean particle size of less than about 20 microns. The host material comprises one or more materials selected from the group consisting of carbonaceous materials, Si, Sn, tin oxides, composite tin alloys, transition metal oxides, lithium metal nitrides and lithium metal oxides. Preferably, the host material in the negative electrode comprises a carbonaceous material and, more preferably, comprises graphite. The amount of lithium metal present in the negative electrode is preferably no more than the maximum amount sufficient to intercalate in, alloy with, or be absorbed by the host material in the negative electrode. For example, if the host material is carbon, the amount of lithium is preferably no more than the amount needed to make LiC_6 .

The present invention also includes a method of preparing a freshly prepared anode for a secondary battery that includes the steps of providing a host material that is capable of absorbing and desorbing lithium in an electrochemical system, dispersing lithium metal in the host material and forming the host material and the lithium metal dispersed therein into an anode. The lithium metal and the host material is preferably mixed together with a non-aqueous liquid to produce a slurry and then applied to a current collector and dried to form the anode. Alternatively, the anode can be formed by chemical means by immersing the host material in a suspension of lithium metal in a non-aqueous liquid, and then formed into an anode.

The present invention further includes a method of operating a secondary battery. First, a freshly prepared, secondary battery is provided that includes a positive electrode including an active material, a negative electrode comprising a host material capable of absorbing and desorbing lithium in an electrochemical system and lithium metal dispersed in the host material, a separator for separating the positive electrode and the negative electrode, and an electrolyte in communication with the positive electrode and the negative electrode. In particular, the secondary battery is manufactured with lithium metal dispersed in the host material of the anode. The freshly assembled battery is in a charged state and more preferably is in a fully charged state (with all the removable lithium present in the anode of the freshly prepared battery). The freshly prepared secondary battery is initially discharged by transmitting lithium ions from the negative electrode to the positive electrode through the electrolyte. The secondary battery can then be charged or recharged by transmitting lithium ions from the positive electrode to the negative electrode through the electrolyte and then discharged again by transmitting lithium ions from the negative electrode to the positive electrode through the electrolyte. The charging and discharging steps can occur for numerous cycles while maintaining the high specific capacities of the cathode active materials and maintaining safe operating conditions.

These and other features and advantages of the present invention will become more readily apparent to those skilled in the art upon consideration of the following detailed description and accompanying drawing, which describe both the preferred and alternative embodiments of the present invention.

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Brief Description of the Drawing

Fig. 1 illustrates a simplified secondary battery construction including a cathode, anode, separator and electrolyte, in accordance with the invention.

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Detailed Description of the Preferred Embodiments of the Invention

In the drawings and the following detailed description, preferred embodiments are described in detail to enable practice of the invention. Although the

invention is described with reference to these specific preferred embodiments, it will be understood that the invention is not limited to these preferred embodiments. But to the contrary, the invention includes numerous alternatives, modifications and equivalents as will become apparent from consideration of the following detailed 5 description and accompanying drawing.

As illustrated in Figure 1, the present invention is a secondary battery 10 that comprises a positive electrode or cathode 12, a negative electrode or anode 14, a separator 16 for separating the positive electrode and the negative electrode, and an electrolyte in electrochemical communication with the positive electrode and the 10 negative electrode. The secondary battery 10 also includes a current collector 20 that is in electrical contact with the cathode and a current collector 22 that is in electrical contact with the anode. The current collectors 20 and 22 are in electrical contact with one another through an external circuit (not shown). The secondary battery 10 can have any construction known in the art such as a "jelly roll" or stacked construction.

15 The cathode 12 is formed of an active material, which is typically combined with a carbonaceous material and a binder polymer. The active material used in the cathode 12 is preferably a material that can be lithiated at a useful voltage (e.g. 2.0 to 5.0 V versus lithium). Preferably, non-lithiated materials such as MnO_2 , V_2O_5 or MoS_2 , or mixtures thereof, can be used as the active material, and more 20 preferably, MnO_2 is used. However, lithiated materials such as $LiMn_2O_4$ that can be further lithiated can also be used. The non-lithiated active materials are preferred because they generally have higher specific capacities than the lithiated active materials in this construction and thus can provide increased power over secondary batteries that include lithiated active materials. Furthermore, because the anode 14 25 includes lithium as discussed below, it is not necessary that the cathode 12 include a lithiated material for the secondary battery 10 to operate. The amount of active material provided in the cathode 12 is preferably sufficient to accept the removable lithium metal present in the anode 14. For example, if MnO_2 is the cathode active material, then one mole of MnO_2 is preferably present in the cathode 12 per mole of 30 lithium in the anode 14 to produce $LiMnO_2$ in the cathode upon discharge.

When cathode active materials are used that can be lithiated such as those described above, the removable lithium that is cycled in the battery is fully provided by the anode 14 and the battery is assembled or prepared in a fully charged state, as is preferred. Nevertheless, the cathode 12 can also include a minor amount of 5 one or more lithiated active materials (e.g. LiCoO_2 or LiNiO_2) that do not further absorb lithium at a voltage between 2.0V and 5.0V and the battery can still be provided in a primarily charged state. In this event, the cathode preferably has less than 10% (molar) and more preferably less than 10% (molar) of the lithiated material (e.g. LiCoO_2 or LiNiO_2) as the active material. Because LiCoO_2 and LiNiO_2 do not 10 further absorb lithium, the presence of these materials in the cathode 12 does not reduce the amount of cathode active material needed to accept the removable lithium from the anode 14.

The anode 14 is formed of a host material 24 capable of absorbing and desorbing lithium in an electrochemical system with lithium metal 26 dispersed in the 15 host material. For example, the lithium present in the anode 14 can intercalate in, alloy with or be absorbed by the host material when the battery (and particularly the anode) is recharged. The host material includes materials capable of absorbing and desorbing lithium in an electrochemical system such as carbonaceous materials; materials containing Si, Sn, tin oxides or composite tin alloys; transition metal oxides 20 such as CoO ; lithium metal nitrides such as $\text{Li}_{3-x}\text{Co}_x\text{N}$ where $0 < x < 0.5$, and lithium metal oxides such as $\text{Li}_4\text{Ti}_5\text{O}_{12}$. The lithium metal 26 is preferably provided in the anode 14 as a finely divided lithium powder. In addition, the lithium metal 26 preferably has a mean particle size of less than about 20 microns, more preferably less than about 10 microns. The lithium metal can be provided as a pyrophoric powder or 25 as a stabilized low pyrophorosity powder, e.g., by treating the lithium metal powder with CO_2 .

The anode 14 is typically capable of reversibly lithiating and delithiating at an electrochemical potential relative to lithium metal of from greater than 0.0 V to less than or equal to 1.5. If the electrochemical potential is 0.0 or less 30 versus lithium, then the lithium metal will not reenter the anode 14 during charging. Alternatively, if the electrochemical potential is greater than 1.5 V versus lithium then

the battery voltage will be undesirably low. Preferably, the amount of lithium metal 26 present in the anode 14 is no more than the maximum amount sufficient to intercalate in, alloy with, or be absorbed by the host material in the anode 14 when the battery is recharged. For example, if the host material 24 is carbon, the amount of lithium 26 is preferably no more than the amount sufficient to make LiC_6 . In other words, the molar ratio of lithium to carbon in the anode is preferably no more than 1:6.

In accordance with the invention, the anode 14 can be prepared by providing a host material that is capable of absorbing and desorbing lithium in an electrochemical system, dispersing lithium metal in the host material, and forming the host material and the lithium metal dispersed therein into an anode. Preferably, the lithium metal and the host material are mixed with a non-aqueous liquid such as tetrahydrofuran (THF) and a binder, and formed into a slurry. The slurry is then used to form the anode 14, for example, by coating the current collector 22 with the slurry and then drying the slurry. The lithium metal can also be provided in the anode by immersing the host material in a suspension containing lithium metal in a non-aqueous liquid such a hydrocarbon solvent (e.g. hexane). The lithium metal used in the suspension is preferably a finely divided lithium powder as discussed above. The host material can be formed into the shape of the anode and then dipped into the lithium metal suspension or it can be combined with the lithium metal suspension to form a slurry and then applied to the current collector and dried to form the anode. The non-aqueous liquid used to form the suspension can be removed by drying the anode (e.g. at an elevated temperature). No matter what method is used, the lithium metal is preferably distributed as well as possible into the host material. Accordingly, as discussed above, the lithium metal 26 preferably has a mean particle size of less than about 20 microns, more preferably less than about 10 microns.

The host material 24 in the anode 14 can include one or more materials capable of absorbing and desorbing lithium in an electrochemical system such as carbonaceous materials; materials containing Si, Sn, tin oxides or composite tin alloys; transition metal oxides such as CoO; lithium metal nitrides such as $\text{Li}_{3-x}\text{Co}_x\text{N}$ where $0 < x < 0.5$; and lithium metal oxides such as $\text{Li}_4\text{Ti}_5\text{O}_{12}$. Preferably, as mentioned

above, the host material 24 preferably includes graphite. In addition, the host material 24 preferably includes a small amount of carbon black (e.g. less than 5% by weight) as a conducting agent.

As shown in Figure 1, the cathode 12 is separated from the anode 14 by 5 an electronic insulating separator 16. Typically, the separator 16 is formed of a material such as polyethylene, polypropylene, or polyvinylidene fluoride (PVDF).

The secondary battery 10 further includes an electrolyte that is in 10 electrochemical communication with the cathode 12 and anode 14. The electrolyte can be non-aqueous liquid, gel or solid and preferably comprises a lithium salt, e.g., LiPF₆. The electrolyte is provided throughout the battery 10 and particularly within the cathode 12, anode 14 and separator 16. Typically, the electrolyte is a liquid, and the cathode 12, anode 14 and separator 16 are porous materials that are soaked in the electrolyte to provide electrochemical communication between these components.

As mentioned above, the battery 10 includes current collectors 20 and 15 22, which are used to transmit electrons to an external circuit. Preferably, the current collector 20 is made of aluminum foil and current collector 22 is made of copper foil.

The battery 10 of the invention can be prepared by methods known in the art and preferably has a layer thickness within the following ranges (from left to right in Figure 1):

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<u>Layer</u>	<u>thickness</u>
Current collector (20)	20-40 μm
Cathode (12)	70-100 μm
Separator (16)	25-35 μm
Anode (14)	70-100 μm
Current collector (22)	20-40 μm

The battery 10 also includes an electrolyte dispersed throughout the cathode 12, anode 14 and separator 16, and a casing (not shown).

In operation, the freshly prepared secondary battery 10 is initially in a 25 charged state, more preferably a fully charged state, and is initially discharged by transmitting lithium ions from the anode 14 to the cathode 12 through the electrolyte. At the same time, electrons are transmitted from the anode 14 to the cathode 12 through the current collector 22, the external circuit, and the current collector 20. The

secondary battery 10 can then be charged or recharged by transmitting lithium ions from the cathode 12 to the anode 14 through the electrolyte and then discharged again as discussed above. The charging and discharging steps can occur for numerous cycles while maintaining the high specific capacities of the cathode active materials and maintaining safe operating conditions.

The secondary battery 10 can be used for various types of applications. For example, the secondary battery can be used in portable electronics such as cellular phones, camcorders, and laptop computers, and in large power applications such as for electric vehicles and hybrid electric vehicles.

10 The present invention provides secondary batteries having a high specific capacity, safe operating conditions and good cycleability. In particular, because lithium metal is provided in the anode, non-lithiated materials can be used as the preferred cathode active material in the secondary battery. These non-lithiated materials have higher specific capacities than the lithiated materials presently used in
15 lithium-ion batteries. Unlike traditional lithium secondary batteries having non-lithiated cathode active materials and metallic lithium anodes, it has been discovered that secondary batteries produced using non-lithiated cathode active materials combined with the anodes of the invention operate safely and do not generate lithium dendrites upon cycling. Furthermore, the secondary batteries of the present invention
20 are safer to operate than lithium-ion batteries, which become unstable when lithium is removed from the cathode during charging. In particular, because the cathode active material in the secondary batteries of the invention is typically in a fully charged state when the battery is freshly prepared, it is more stable than the cathode materials used in lithium-ion batteries. Moreover, the batteries of the invention can be charged and
25 discharged numerous times while maintaining safe operating conditions and the high specific capacities of the cathode active materials.

It is understood that upon reading the above description of the present invention and reviewing the accompanying drawings, one skilled in the art could make changes and variations therefrom. These changes and variations are included in
30 the spirit and scope of the following appended claims.

CLAIMS

1. A freshly prepared anode for a secondary battery comprising a host material that is capable of absorbing and desorbing lithium in an electrochemical system and lithium metal dispersed in the host material.
2. An anode as claimed in claim 1, wherein the lithium metal is a finely divided lithium powder.
3. An anode as claimed in claim 2, wherein the lithium powder has a mean particle size of less than about 20 microns.
4. An anode as claimed in any one of the preceding claims, being capable of reversibly lithiating and delithiating at an electrochemical potential relative to lithium metal of from greater than 0.0V to less than or equal to 1.5 V.
5. An anode as claimed in any one of the preceding claims, wherein the host material comprises one or more materials capable of reversibly lithiating and delithiating at an electrochemical potential versus lithium of from greater than 0.0 V to less than or equal to 1.5 V.
6. An anode as claimed in any one of the preceding claims, wherein the host material comprises one or more materials selected from the group consisting of carbonaceous materials, Si, Sn, tin oxides, composite tin alloys, transition metal oxides, lithium metal nitrides and lithium metal oxides.
7. An anode as claimed in any one of the preceding claims, wherein the host material comprises a carbonaceous material.
8. An anode as claimed in claim 7, wherein the carbonaceous material is

graphite.

9. An anode as claimed in claim 8, wherein the host material further comprises carbon black.

10. An anode as claimed in any one of the preceding claims, wherein the amount of lithium metal in said anode is no more than the maximum amount sufficient to intercalate in, alloy with, or be absorbed by the host material in said anode when the anode is recharged.

11. A secondary battery including an anode as claimed in any one of the preceding claims.

12. A freshly prepared, secondary battery comprising:

a positive electrode including an active material;

a negative electrode comprising a host material that is capable of absorbing and desorbing lithium in an electrochemical system and lithium metal dispersed in the host material;

a separator for separating the positive electrode and the negative electrode;

and

an electrolyte in communication with the positive electrode and the negative electrode.

13. A secondary battery as claimed in claim 12, wherein the positive electrode includes a compound that can be lithiated at an electrochemical potential of 2.0 to 5.0V versus lithium as the active material.

14. A secondary battery as claimed in claim 12 or 13, wherein the active

material in the positive electrode is selected from the group consisting of MnO_2 , V_2O_5 and MoS_2 , and mixtures thereof.

15. A secondary battery as claimed in any one of claims 12 to 14, wherein the active material in the positive electrode comprises MnO_2 .

16. A secondary battery as claimed in any one of claims 12 to 14, wherein the active material in the positive electrode is LiMn_2O_4 .

17. A secondary battery as claimed in any one of claims 12 to 15, wherein the lithium metal in said anode is a finely divided lithium powder.

18. A secondary battery as claimed in claim 17, wherein the lithium metal in said anode has a mean particle size of less than about 20 microns.

19. A secondary battery as claimed in any one of claims 12 to 18, wherein the anode is capable of reversibly lithiating and delithiating at an electrochemical potential relative to lithium metal of from greater than 0.0 V to less than or equal to 1.5 V.

20. A secondary battery as claimed in any one of claims 12 to 19, wherein the host material in said anode comprises one or more materials capable of reversibly lithiating and delithiating at an electrochemical potential versus lithium of from greater than 0.0 V to less than or equal to 1.5 V.

21. A secondary battery as claimed in any one of claims 12 to 20, wherein the host material in said anode comprises one or more materials selected from the group consisting of carbonaceous materials, Si, Sn, tin oxides, composite tin alloys, transition metal oxides, lithium metal nitrides and lithium metal oxides.

22. A secondary battery as claimed in any one of claims 12 to 21, wherein the host material in said anode comprises a carbonaceous material.

23. A secondary battery as claimed in claim 22, wherein the carbonaceous material is graphite.

24. A secondary battery as claimed in claim 23, wherein the host material further comprises carbon black.

25. A secondary battery as claimed in any one of claims 12 to 24, wherein the amount of lithium metal in said negative electrode is no more than the maximum amount sufficient to intercalate in, alloy with, or be absorbed by the host material in said anode when said battery is recharged.

26. A secondary battery as claimed in any one of claims 12 to 25, wherein the amount of active material in said cathode is sufficient to accept the removable lithium metal present in said anode.

27. A secondary battery as claimed in any one of claims 12 to 26, being in a fully charged state.

28. A method of preparing an anode for a secondary battery comprising:
providing a host material that is capable of absorbing and desorbing lithium in an electrochemical system;
dispersing lithium metal in the host material; and
forming the host material and the lithium metal dispersed therein into an anode.

29. A method as claimed in claim 28, wherein said dispersing step comprises

mixing the lithium metal, the host material and a non-aqueous liquid together to form a slurry.

30. A method as claimed in claim 29, wherein said forming step comprising applying the slurry to a current collector and drying the slurry.

31. A method as claimed in any one of claims 28 to 30, wherein said dispersing step comprises immersing the host material in a suspension containing lithium metal and a non-aqueous liquid.

32. A method as claimed claim 31, wherein said dispersing step comprises immersing the host material in a suspension of lithium metal in a hydrocarbon.

33. A method as claimed in any one of claims 28 to 32, wherein said dispersing step comprises dispersing a finely divided lithium metal powder in the host material.

34. A method as claimed in claim 33, wherein said dispersing step comprises dispersing lithium metal having a mean particle size of less than about 20 microns in the host material.

35. A method as claimed in any one of claims 28 to 34, wherein said providing step includes providing a host material comprising one or more materials capable of reversibly lithiating and delithiating at an electrochemical potential versus lithium of from greater than 0.0 V to less than or equal to 1.5 V.

36. A method as claimed in any one of claims 28 to 35, wherein said providing step comprises providing a host material comprising one or more materials selected from the group consisting of carbonaceous materials, Si, Sn, tin oxides,

composite tin alloys, transition metal oxides, lithium metal nitrides, and lithium metal oxides.

37. A method as claimed in any one of claims 28 to 36, wherein said providing step comprises providing a host material comprising a carbonaceous material.

38. A method as claimed in claim 37, wherein said providing step comprises providing a host material wherein the carbonaceous material is graphite.

39. A method as claimed in claim 38, wherein said providing step comprises providing a host material wherein the host material further comprises carbon black.

40. A method of operating a secondary battery comprising the steps of:

(a) providing a freshly prepared, secondary battery comprising a positive electrode including an active material, a negative electrode comprising a host material capable of absorbing and desorbing lithium in an electrochemical system and lithium metal dispersed in the host material, a separator for separating the positive electrode and the negative electrode, and an electrolyte in communication with the positive electrode and the negative electrode;

(b) initially discharging the secondary battery by transmitting lithium ions from the negative electrode to the positive electrode through the electrolyte;

(c) charging the secondary battery by transmitting lithium ions from the positive electrode to the negative electrode through the electrolyte;

(d) discharging the secondary battery by transmitting lithium ions from the negative electrode to the positive electrode through the electrolyte;

(e) repeating steps (c) and (d).

41. A freshly prepared anode substantially as hereinbefore described with reference to and as illustrated in the accompanying the drawing.

42. A secondary battery substantially as hereinbefore described with reference to and as illustrated in the accompanying the drawing.

43. A method of preparing an anode substantially as hereinbefore described with reference to and as illustrated in the accompanying the drawing.

44. A method of providing a secondary battery substantially as hereinbefore described with reference to and as illustrated in the accompanying the drawing.



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Claims searched: All claims

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Examiner: A.R.Martin
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Patents Act 1977
Search Report under Section 17

Databases searched:

UK Patent Office collections, including GB, EP, WO & US patent specifications, in:

UK Cl (Ed. T): H1B

Int Cl (Ed. 7): H01M 4/00

Other: On line databases WPI, EPODOC, JAPIO

Documents considered to be relevant:

Category	Identity of document and relevant passage		Relevant to claims
X	US 6270926 A	Murata see claim 1	1, 11, 12, 28, 40 at least
X	US 6156457 A	Toshiba see claim 1	"
X	US 5958622 A	Toyota see abstract	"
X	US 5753387 A	Toshiba see claim 1	"
X	US 5725968 A	Honda see abstract	"
X	US 5312611 A	Toshiba see abstract	"
X	EP 0696075 A	Seiko see abstract	"
X	EP 0647974 A	Toshiba see abstract	"
X	EP 0601832 A	Honda see abstract	"
X	EP 0573266 A	Toshiba see abstract	"
X	EP 0763865 A	Toshiba see abstract	"

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Y	Document indicating lack of inventive step if combined with one or more other documents of same category.	P	Document published on or after the declared priority date but before the filing date of this invention.
&	Member of the same patent family	E	Patent document published on or after, but with priority date earlier than, the filing date of this application.